

## AMENDMENT

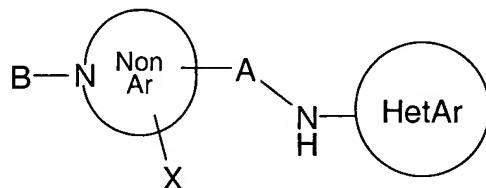
Please amend the application, without prejudice, without admission, without surrender of subject matter, and without any intention of creating any estoppel as to equivalents, as follows:

### Amendment to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

### Listing of Claims:

Claim 1(currently amended): A compound having the formula (I):



(I)

or a pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic [5-7] 6 membered ring containing 1 [or 2] nitrogen ring atoms or an aza-bicyclo-octane ring;

HetAr is a 5 or 6 membered heteroaromatic ring containing 1-3 nitrogen ring atoms, or isoxazolyl, thiazolyl, thiadiazolyl, quinolinyl, quinazolinyl, purinyl, pteridinyl, benzimidazolyl, pyrrolopyrimidinyl, or imidazopyridinyl;

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>1</sub>-4alkoxy, C<sub>2</sub>-4alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1</sub>-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-;

A is  $-C[[0]]\underline{1-4alkyl}-$ ;

B is  $aryl(CH_2)_{0-3}-O-C(O)-$ , heteroaryl( $CH_2)_{1-3}-O-C(O)-$ , indanyl( $CH_2)_{0-3}-O-C(O)-$ ,  $aryl(CH_2)_{1-3}-C(O)-$ , aryl-cyclopropyl-C(O)-, heteroaryl-cyclopropyl-C(O)-, heteroaryl( $CH_2)_{1-3}-C(O)-$ ,  $[[aryl(CH_2)_{1-3}-]]$ ,  $[[heteroaryl(CH_2)_{1-3}-]]$  aryl( $CH_2)_{1-3}-NH-C(O)-$ , aryl( $CH_2)_{1-3}-NH-C(NCN)-$ , aryl( $CH_2)_{1-3}-SO_2-$ , heteroaryl( $CH_2)_{1-3}-SO_2-$ , wherein any of the aryl or heteroaryl is optionally substituted by 1-5 substitutents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>3-6</sub>cycloalkyl, C<sub>1-4</sub>alkoxy, trifluoromethyl, bromo, fluoro, or chloro; and

X is H, OH, F, C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, NH<sub>2</sub>, or X taken with an adjacent bond is =O.

Claim 2(previously presented): The compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

B is  $aryl(CH_2)_{0-3}-O-C(O)-$ , wherein the aryl is optionally substituted by 1-5 substitutents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>3-6</sub>cycloalkyl, C<sub>1-4</sub>alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

Claim 3(previously presented): The compound according to Claim 2, or a pharmaceutically acceptable salt thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 1 nitrogen ring atom;

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-,  $-N(C_0-4alkyl)(C_0-4alkyl)$ , nitro,  $(C_1-2alkyl)(C_1-2alkyl)NCH_2-$ ,  $(C_1-2alkyl)HNCH_2-$ , Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 4(previously presented): The compound according to Claim 2, or a pharmaceutically acceptable salt thereof, wherein

HetAr is an isoxazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-,

phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 5(previously presented): The compound according to Claim 2, or a pharmaceutically acceptable salt thereof, wherein

HetAr is a thiadiazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 6(previously presented): The compound according to Claim 2, or a pharmaceutically acceptable salt thereof, wherein

HetAr is a 5 membered heteroaromatic ring containing 2 nitrogen ring atoms;

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 7(previously presented): The compound according to Claim 2, or a pharmaceutically acceptable salt thereof, wherein

HetAr is quinolinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 8(previously presented): The compound according to Claim 2, or a pharmaceutically acceptable salt thereof, wherein

HetAr is purinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 9(previously presented): The compound according to Claim 2, or a pharmaceutically acceptable salt thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 2 nitrogen ring atoms;

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>1</sub>-4alkoxy, C<sub>2</sub>-4alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1</sub>-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 10(previously presented): The compound according to Claim 2, or a pharmaceutically acceptable salt thereof, wherein

HetAr is thiazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>1</sub>-4alkoxy, C<sub>2</sub>-4alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1</sub>-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 11(previously presented): The compound according to Claim 2, or a pharmaceutically acceptable salt thereof, wherein

HetAr is pteridinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>1</sub>-4alkoxy, C<sub>2</sub>-4alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1</sub>-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 12(previously presented): The compound according to Claim 2, or a pharmaceutically acceptable salt thereof, wherein

HetAr is pyrrolopyrimidinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 13(previously presented): The compound according to Claim 2, or a pharmaceutically acceptable salt thereof, wherein

HetAr is a imidazopyridinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 14(previously presented): The compound according to Claim 2, or a pharmaceutically acceptable salt thereof, wherein

HetAr is benzimidazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 15(previously presented): The compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and B is aryl(CH<sub>2</sub>)<sub>1-3</sub>-SO<sub>2</sub>-, wherein the aryl is optionally substituted by 1-5 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>3-6</sub>cycloalkyl, C<sub>1-4</sub>alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

Claim 16(previously presented): The compound according to Claim 15, or a pharmaceutically acceptable salt thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 2 nitrogen ring atoms;

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>1</sub>-4alkoxy, C<sub>2</sub>-4alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1</sub>-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 17(previously presented): The compound according to Claim 15, or a pharmaceutically acceptable salt thereof, wherein

HetAr is quinazolinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>1</sub>-4alkoxy, C<sub>2</sub>-4alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1</sub>-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 18(previously presented): The compound according to Claim 15, or a pharmaceutically acceptable salt thereof, wherein

HetAr is purinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>1</sub>-4alkoxy, C<sub>2</sub>-4alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1</sub>-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 19(previously presented): The compound according to Claim 15, or a pharmaceutically acceptable salt thereof, wherein

HetAr is imidazopyridinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>1</sub>-4alkoxy, C<sub>2</sub>-4alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1</sub>-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 20(previously presented): The compound according to Claim 15, or a pharmaceutically acceptable salt thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 1 nitrogen ring atom; and HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 21(withdrawn): The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 5 membered ring containing 1 nitrogen ring atom; and B is aryl(CH<sub>2</sub>)<sub>0-3</sub>-O-C(O)-, wherein the aryl is optionally substituted by 1-5 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>3-6</sub>cycloalkyl, C<sub>1-4</sub>alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

Claim 22(withdrawn): The compound according to Claim 21, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 2 nitrogen ring atoms; HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 23(withdrawn): The compound according to Claim 21, or pharmaceutically acceptable salts thereof, wherein

HetAr is pteridinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 24(withdrawn): The compound according to Claim 21, or pharmaceutically acceptable salts thereof, wherein

HetAr is purinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 25(withdrawn): The compound according to Claim 21, or pharmaceutically acceptable salts thereof, wherein

HetAr is benzimidazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 26(withdrawn): The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is an aza bicyclo octane ring; and

B is aryl(CH<sub>2</sub>)<sub>0-3</sub>-O-C(O)-, wherein the aryl is optionally substituted by 1-5 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>3-6</sub>cycloalkyl, C<sub>1-4</sub>alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

Claim 27(withdrawn): The compound according to Claim 26, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 1 nitrogen ring atom; and

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>-, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 28(withdrawn): The compound according to Claim 26, or pharmaceutically acceptable salts thereof, wherein

HetAr is purinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>1</sub>-4alkoxy, C<sub>2</sub>-4alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1</sub>-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 29(withdrawn): The compound according to Claim 26, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 2 nitrogen ring atom; and

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>1</sub>-4alkoxy, C<sub>2</sub>-4alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1</sub>-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 30(withdrawn): The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is an aza bicyclo octane ring; and

B is aryl(CH<sub>2</sub>)<sub>1-3</sub>-SO<sub>2</sub>-, wherein the aryl is optionally substituted by 1-5 substitutents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>3</sub>-6cycloalkyl, C<sub>1</sub>-4alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

Claim 31(previously presented): The compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

B is heteroaryl(CH<sub>2</sub>)<sub>1-3</sub>-C(O)-, wherein the heteroaryl is optionally substituted by 1-5 substitutents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>3</sub>-6cycloalkyl, C<sub>1</sub>-4alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

Claim 32(previously presented): The compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and B is aryl(CH<sub>2</sub>)<sub>1-3</sub>—C(O)—, wherein the aryl is optionally substituted by 1-5 substitutents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>3-6</sub>cycloalkyl, C<sub>1-4</sub>alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

Claim 33(previously presented): The compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and B is aryl—cyclopropyl—C(O)—, wherein the aryl is optionally substituted by 1-5 substitutents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>3-6</sub>cycloalkyl, C<sub>1-4</sub>alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

Claim 34(previously presented): The compound according to Claim 33, or a pharmaceutically acceptable salt thereof, wherein

HetAr is pyridyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl—, phenylethynyl—, heteroarylethynyl—, —N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>—, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>—, Si(CH<sub>3</sub>)<sub>3</sub>—C—, or NH<sub>2</sub>C(O)—.

Claim 35(previously presented): The compound according to Claim 33, or a pharmaceutically acceptable salt thereof, wherein

HetAr is pyrazinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>2-4</sub>alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1-4</sub>alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl—, phenylethynyl—, heteroarylethynyl—, —N(C<sub>0-4</sub>alkyl)(C<sub>0-4</sub>alkyl), nitro, (C<sub>1-2</sub>alkyl)(C<sub>1-2</sub>alkyl)NCH<sub>2</sub>—, (C<sub>1-2</sub>alkyl)HNCH<sub>2</sub>—, Si(CH<sub>3</sub>)<sub>3</sub>—C—, or NH<sub>2</sub>C(O)—.

Claim 36(previously presented): The compound according to Claim 33, or a pharmaceutically acceptable salt thereof, wherein

HetAr is pyridazinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>1</sub>-4alkoxy, C<sub>2</sub>-4alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1</sub>-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 37(previously presented): The compound according to Claim 33, or a pharmaceutically acceptable salt thereof, wherein

HetAr is pyrimidinyl optionally substituted with 1 or 2 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>1</sub>-4alkoxy, C<sub>2</sub>-4alkynyl, trifluoromethyl, hydroxy, hydroxyC<sub>1</sub>-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C<sub>0</sub>-4alkyl)(C<sub>0</sub>-4alkyl), nitro, (C<sub>1</sub>-2alkyl)(C<sub>1</sub>-2alkyl)NCH<sub>2</sub>-, (C<sub>1</sub>-2alkyl)HNCH<sub>2</sub>-, Si(CH<sub>3</sub>)<sub>3</sub>-C-, or NH<sub>2</sub>C(O)-.

Claim 38(previously presented): The compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein

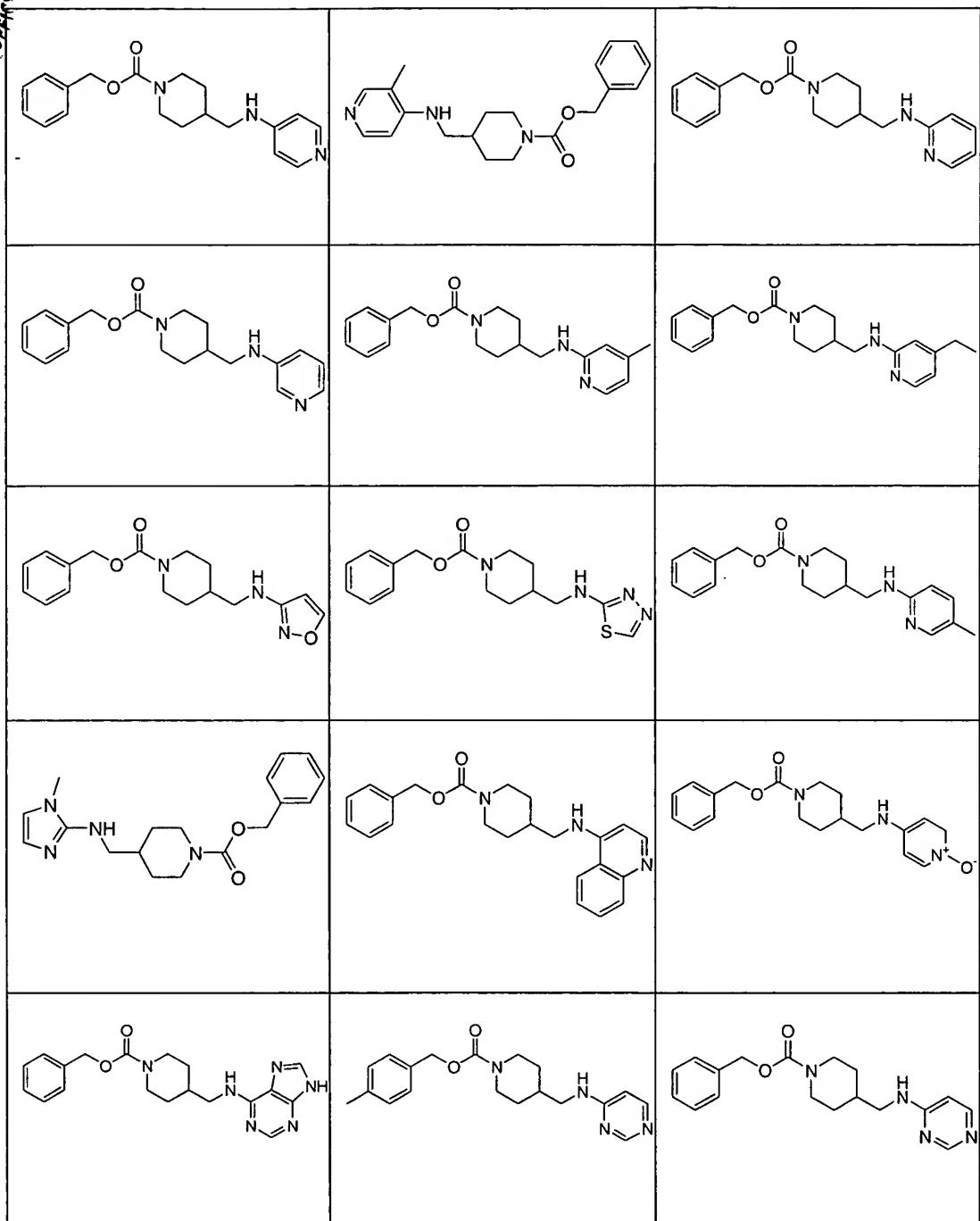
NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and B is heteroaryl(CH<sub>2</sub>)<sub>1-3</sub>-O-C(O)-, wherein the heteroaryl is optionally substituted by 1-5 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>3</sub>-6cycloalkyl, C<sub>1</sub>-4alkoxy, trifluoromethyl, bromo, fluoro, or chloro;.

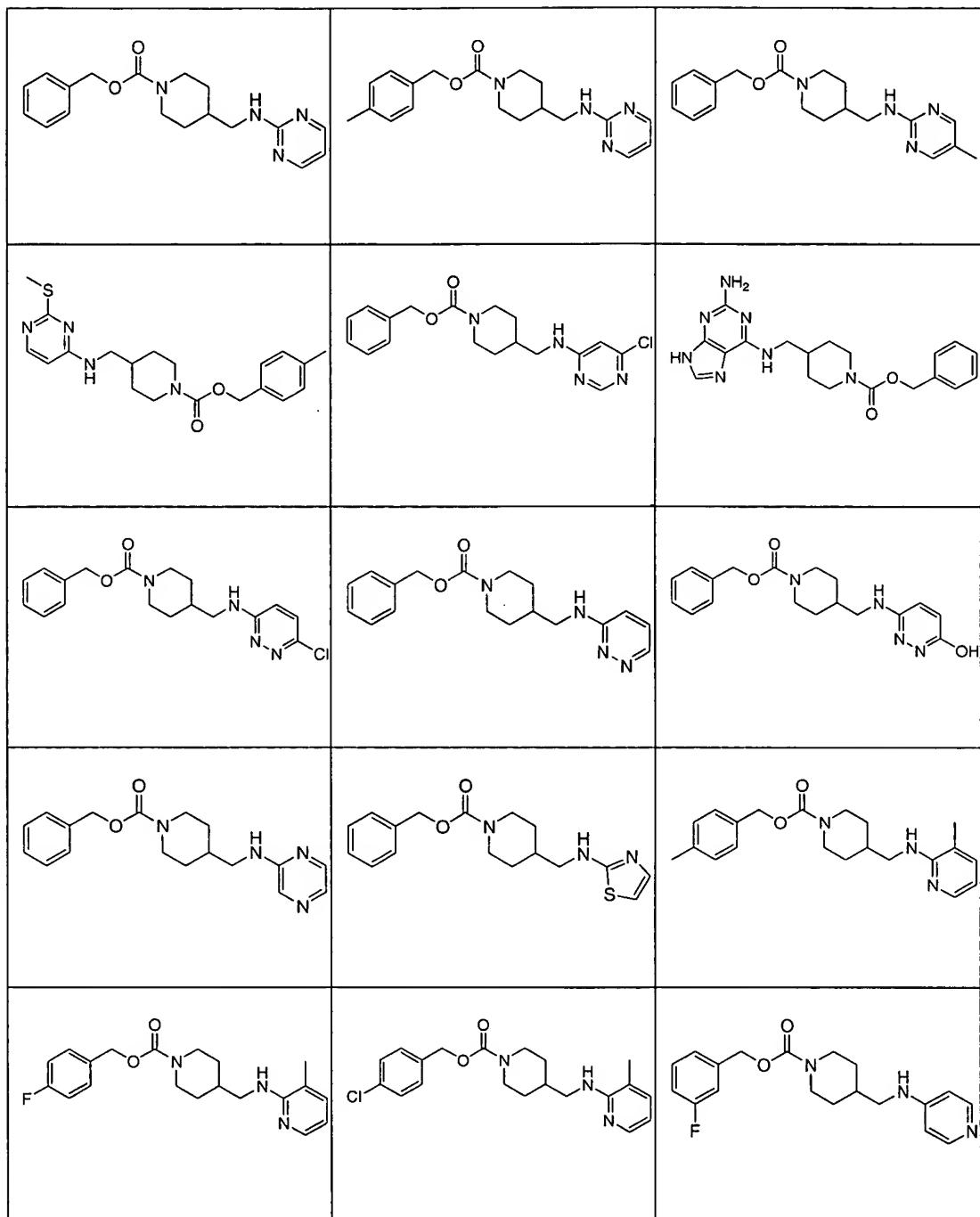
Claim 39(previously presented): The compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein

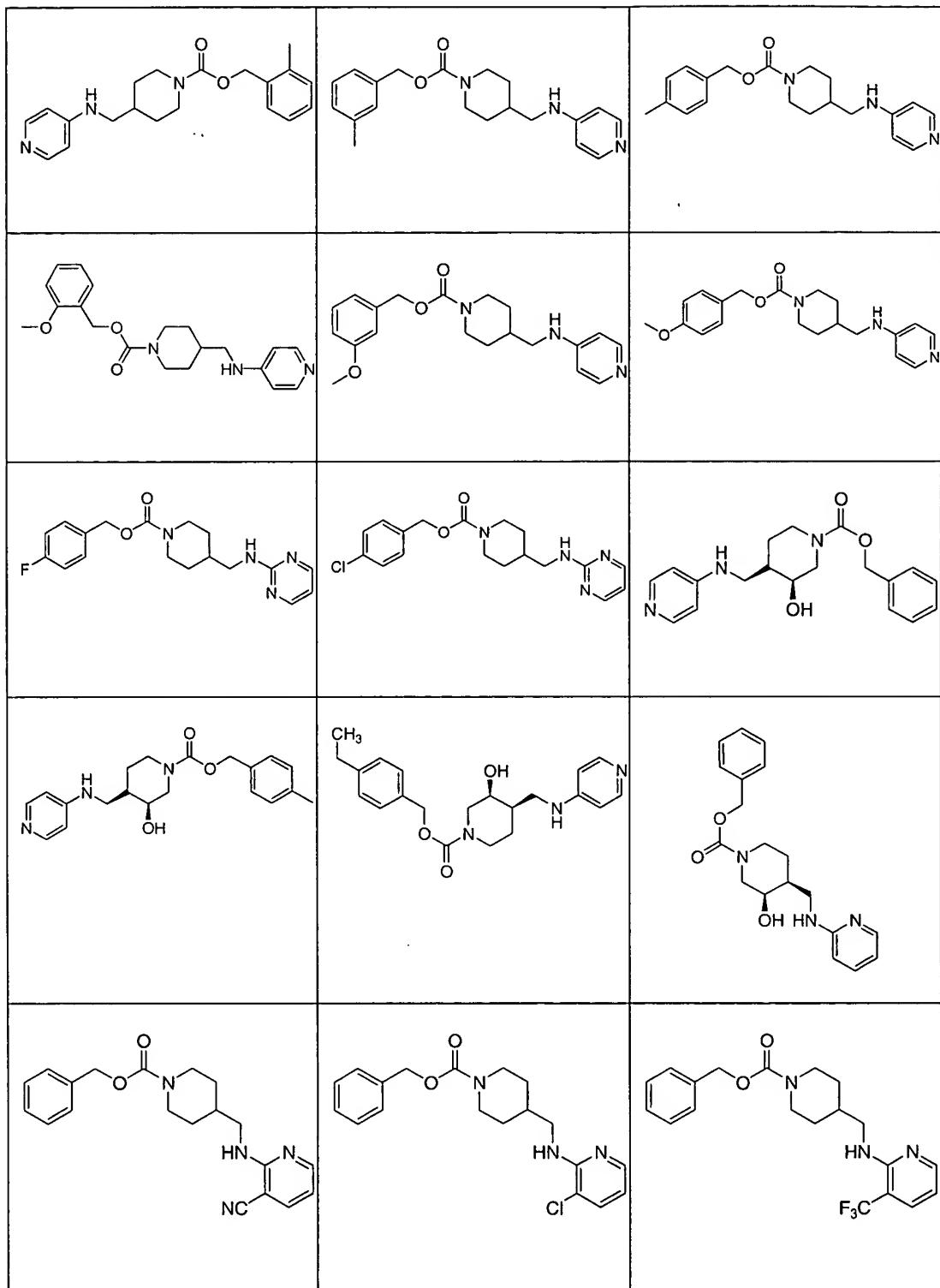
NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and B is aryl(CH<sub>2</sub>)<sub>1-3</sub>-NH-C(NCN)-, wherein the aryl is optionally substituted by 1-5 substituents, each substituent independently is C<sub>1</sub>-4alkyl, C<sub>3</sub>-6cycloalkyl, C<sub>1</sub>-4alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

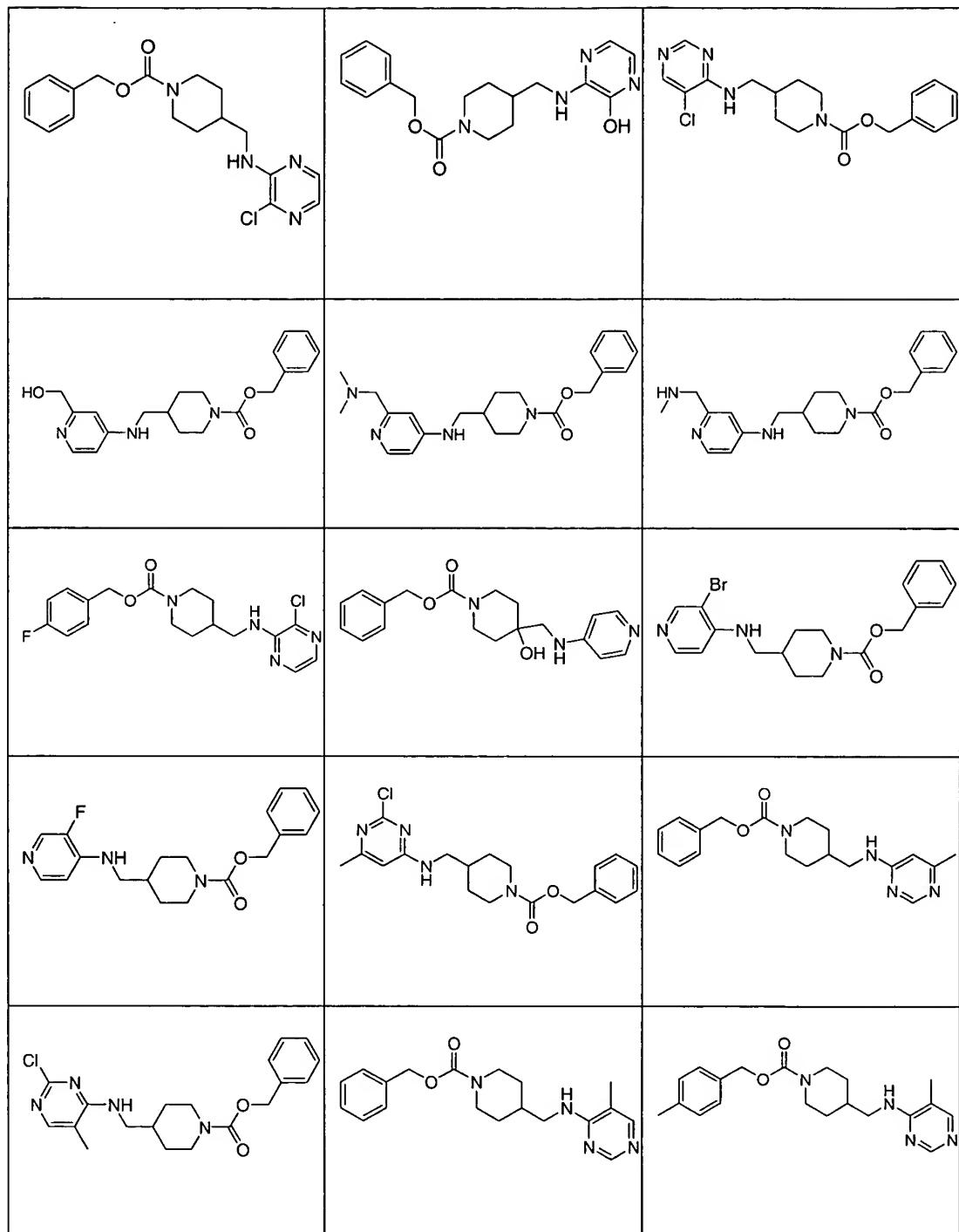
Claim 40(original): The compound according to Claim 1, wherein said compound is

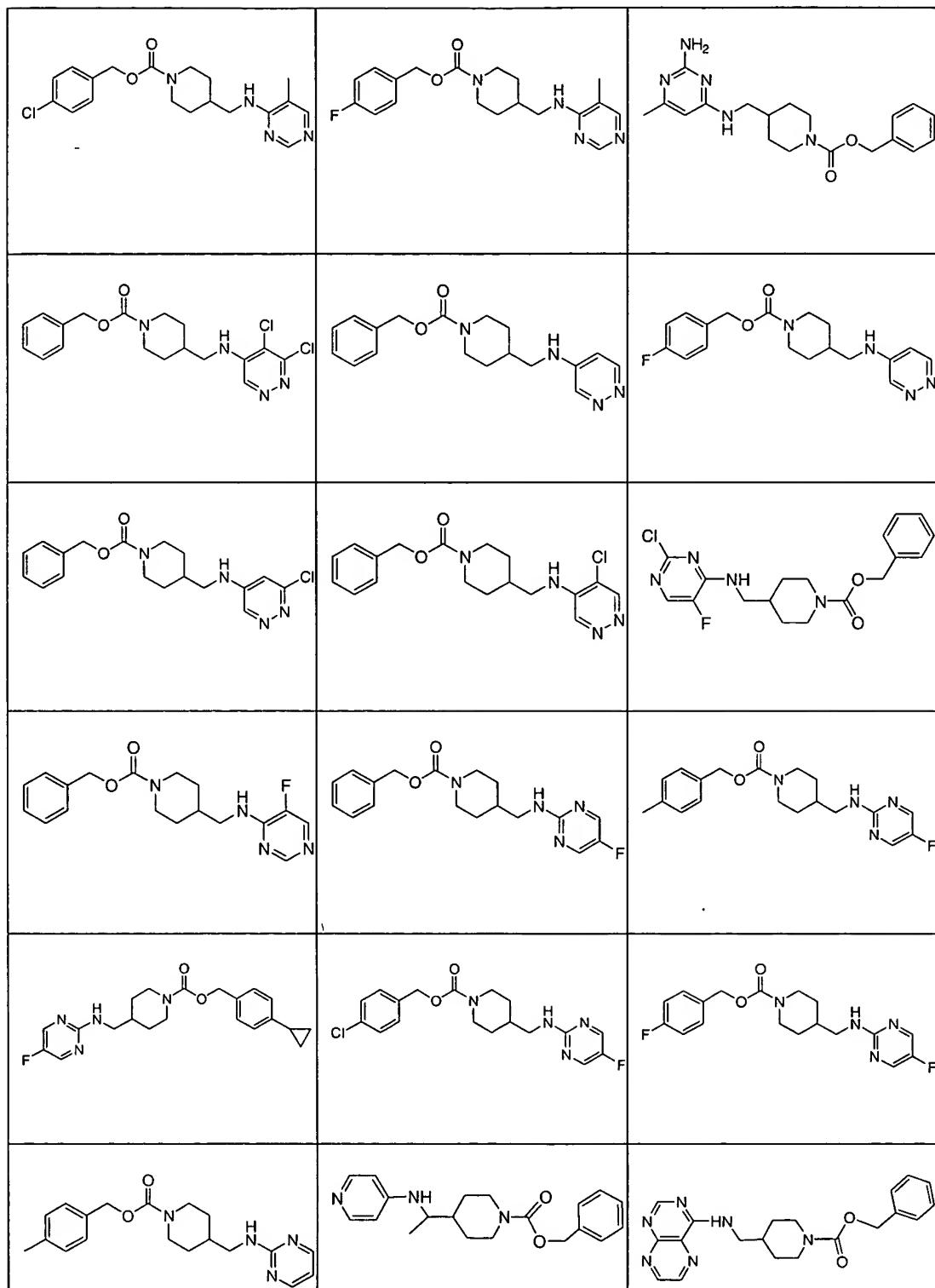
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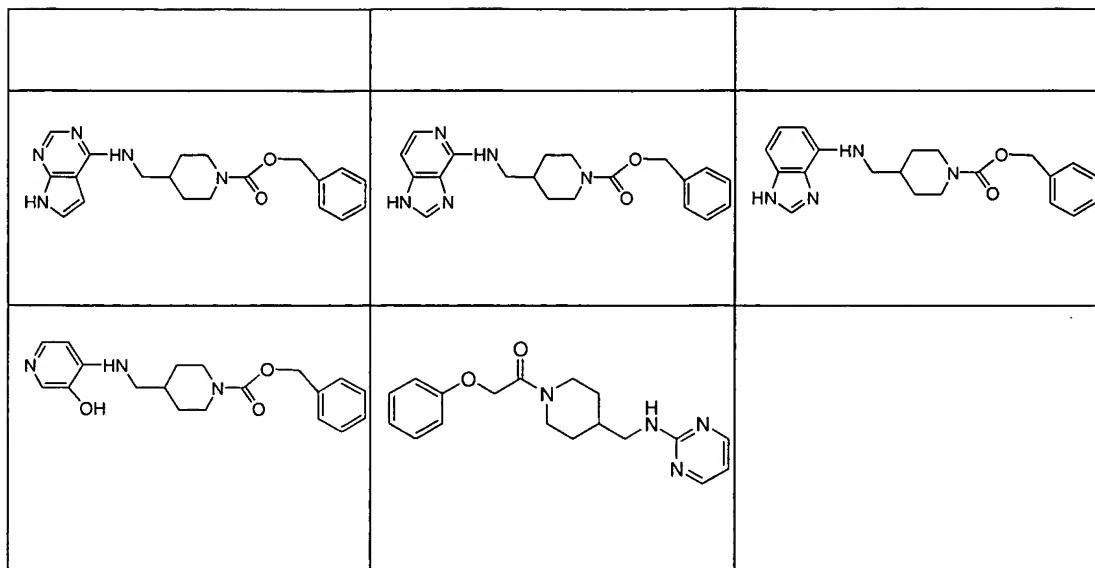






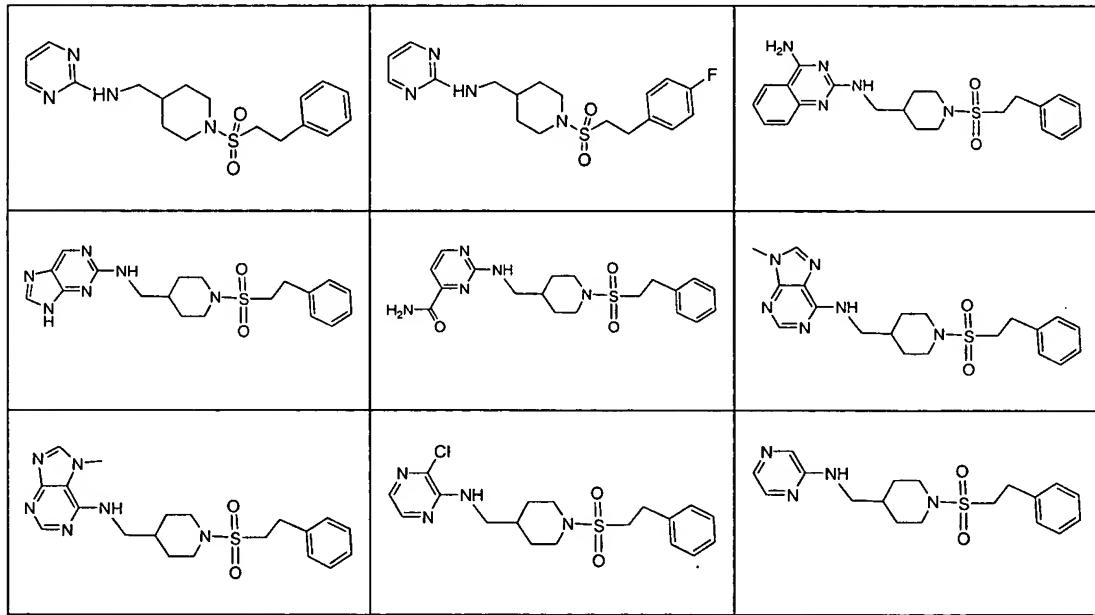


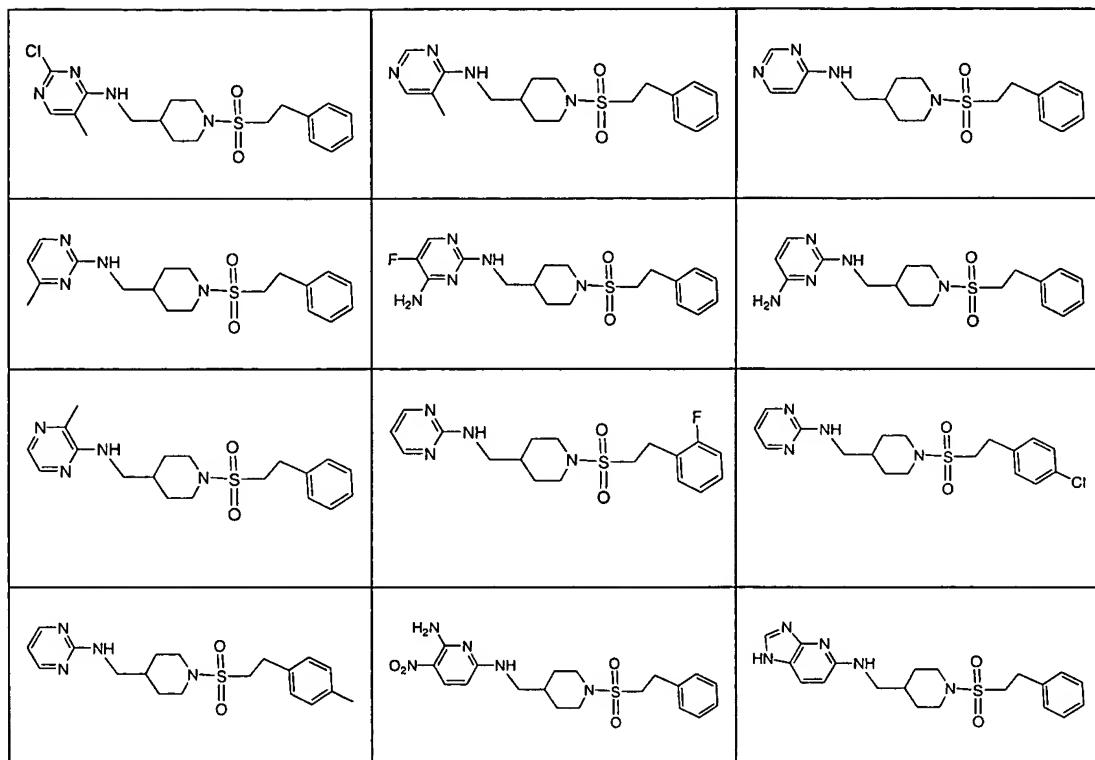




or a pharmaceutically acceptable salt thereof.

Claim 41(original): The compound according to Claim 1, wherein said compound is

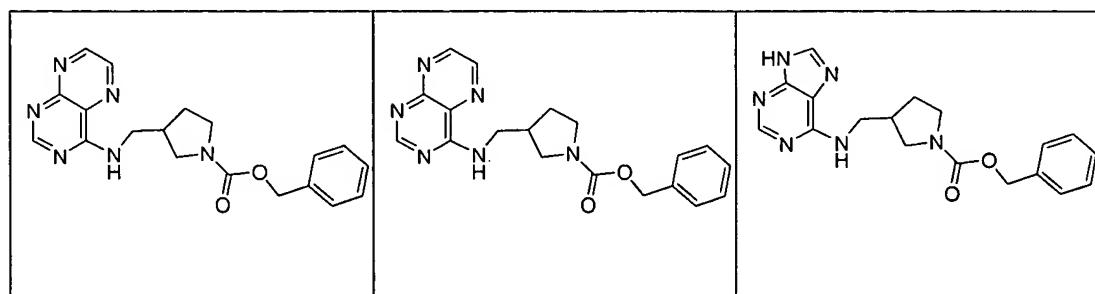




or a pharmaceutically acceptable salt thereof.

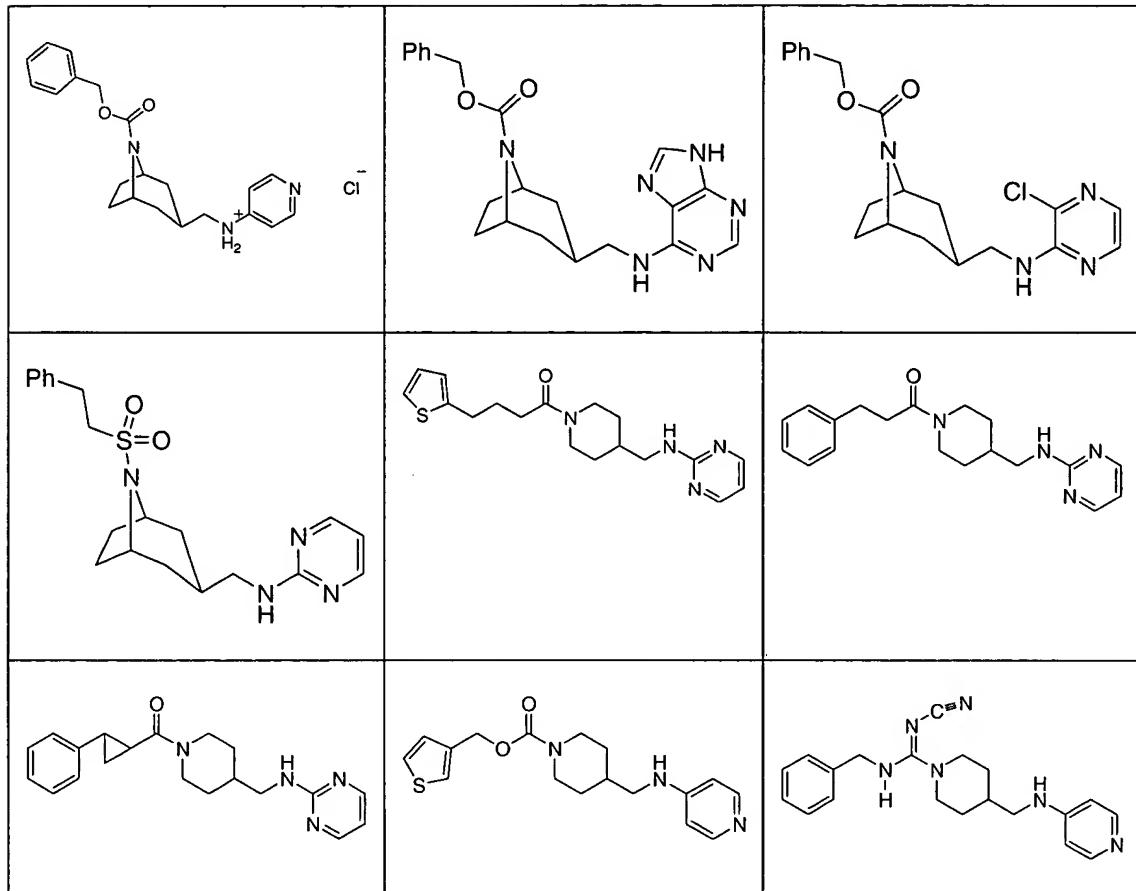
Claim 42(currently amended):  
A compound is represented by

~~The compound according to Claim 1, wherein said~~



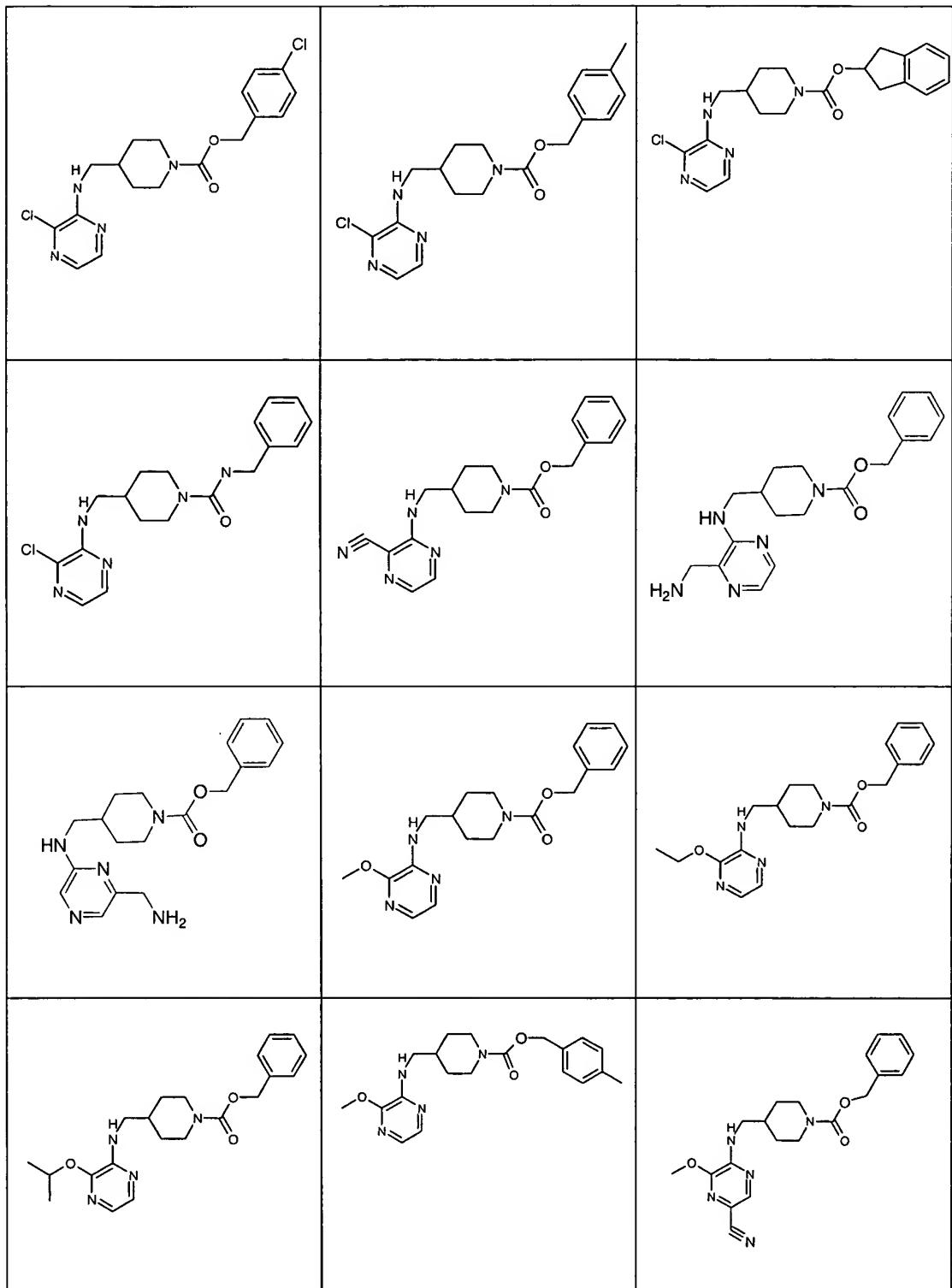
or a pharmaceutically acceptable salt thereof.

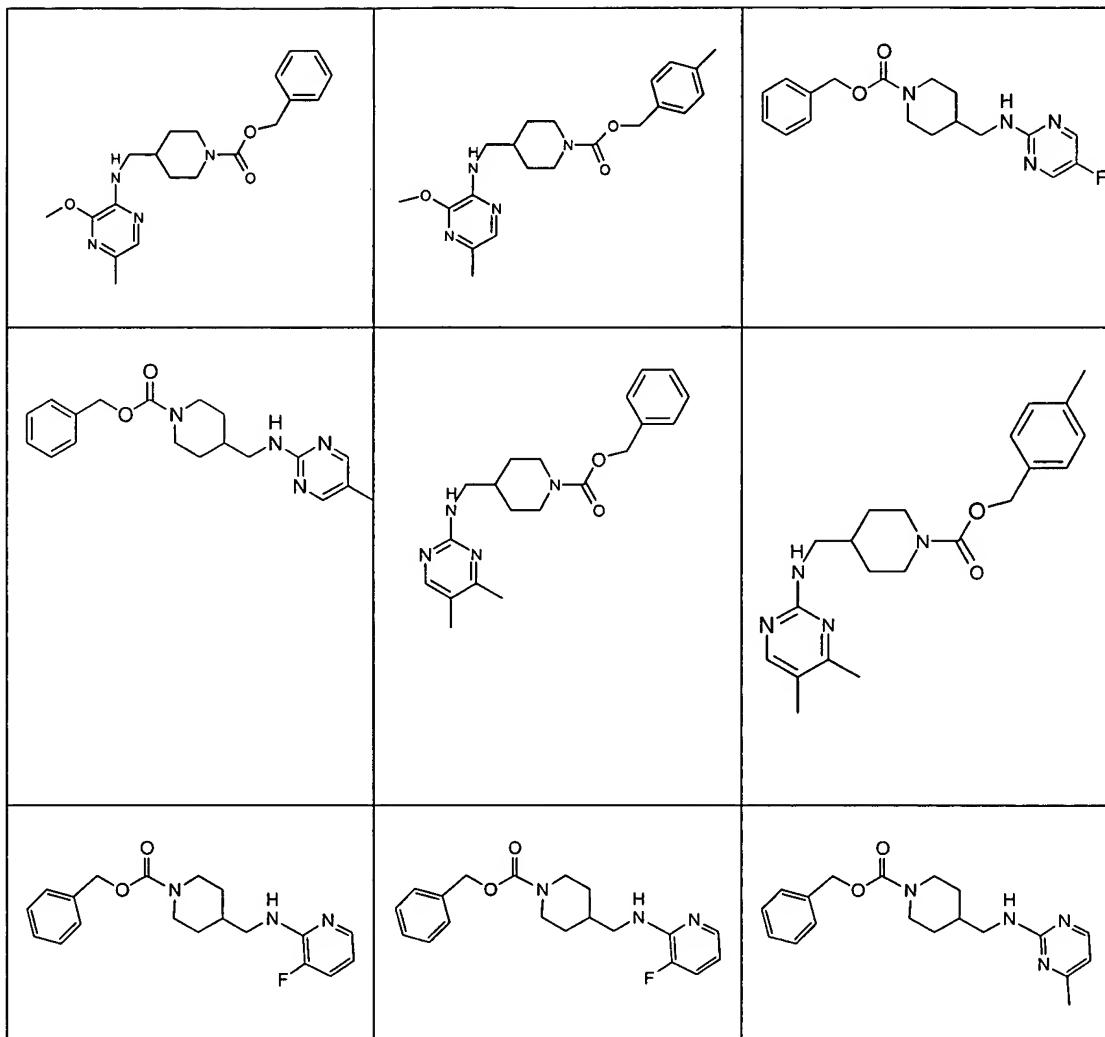
Claim 43 (currently amended): ~~The compound according to Claim 1, wherein said A~~  
compound is represented by



or a pharmaceutically acceptable salt thereof.

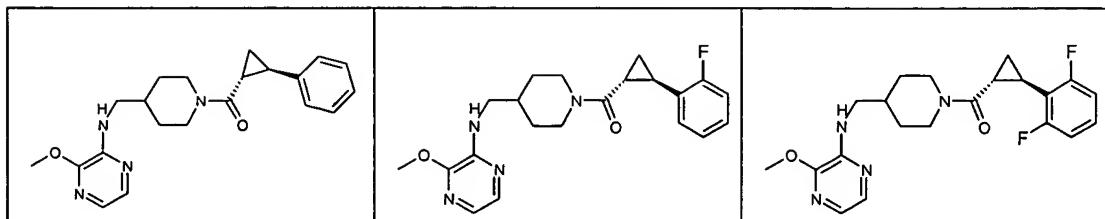
Claim 44 (previously presented): The compound according to Claim 1, wherein said compound is

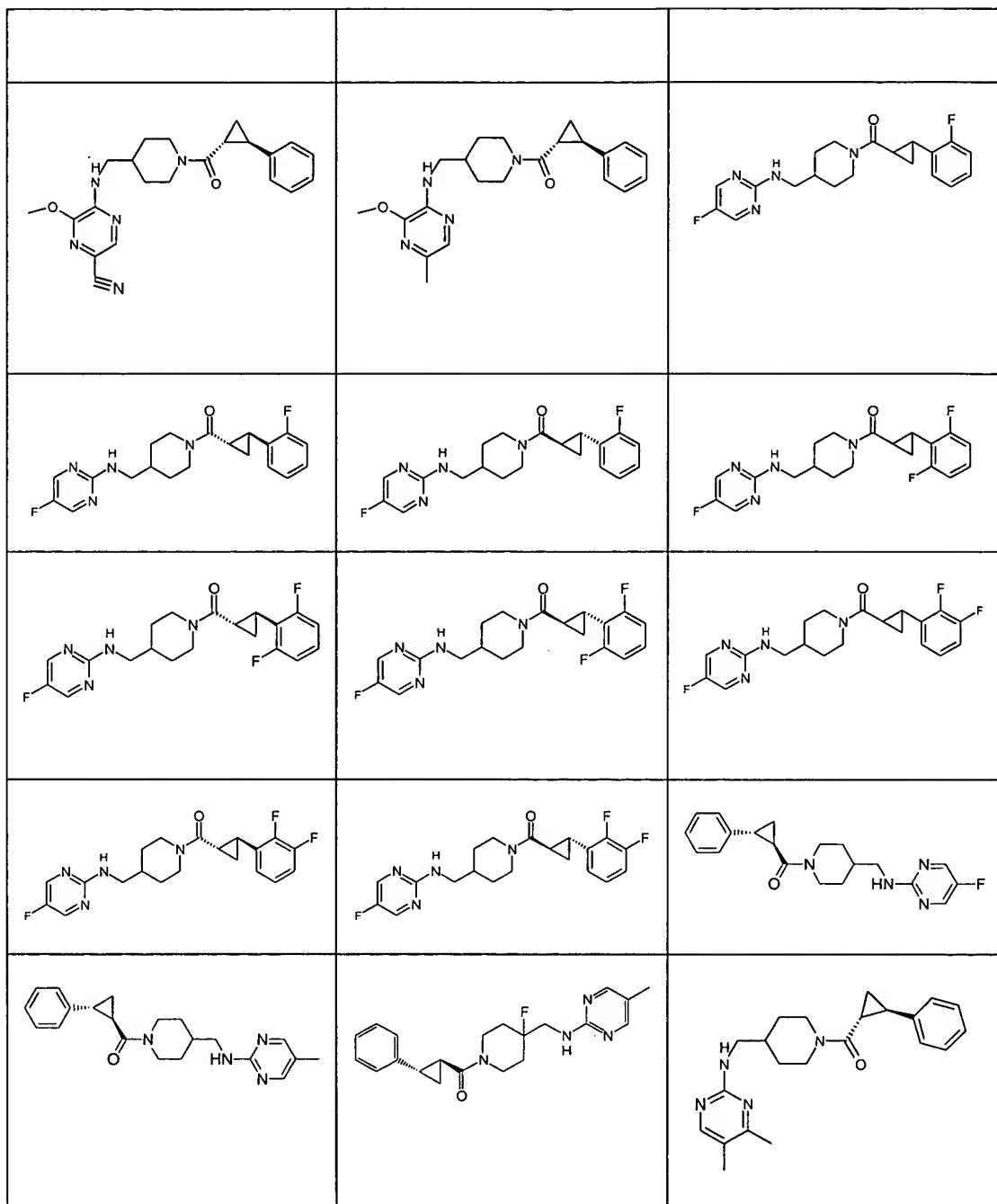


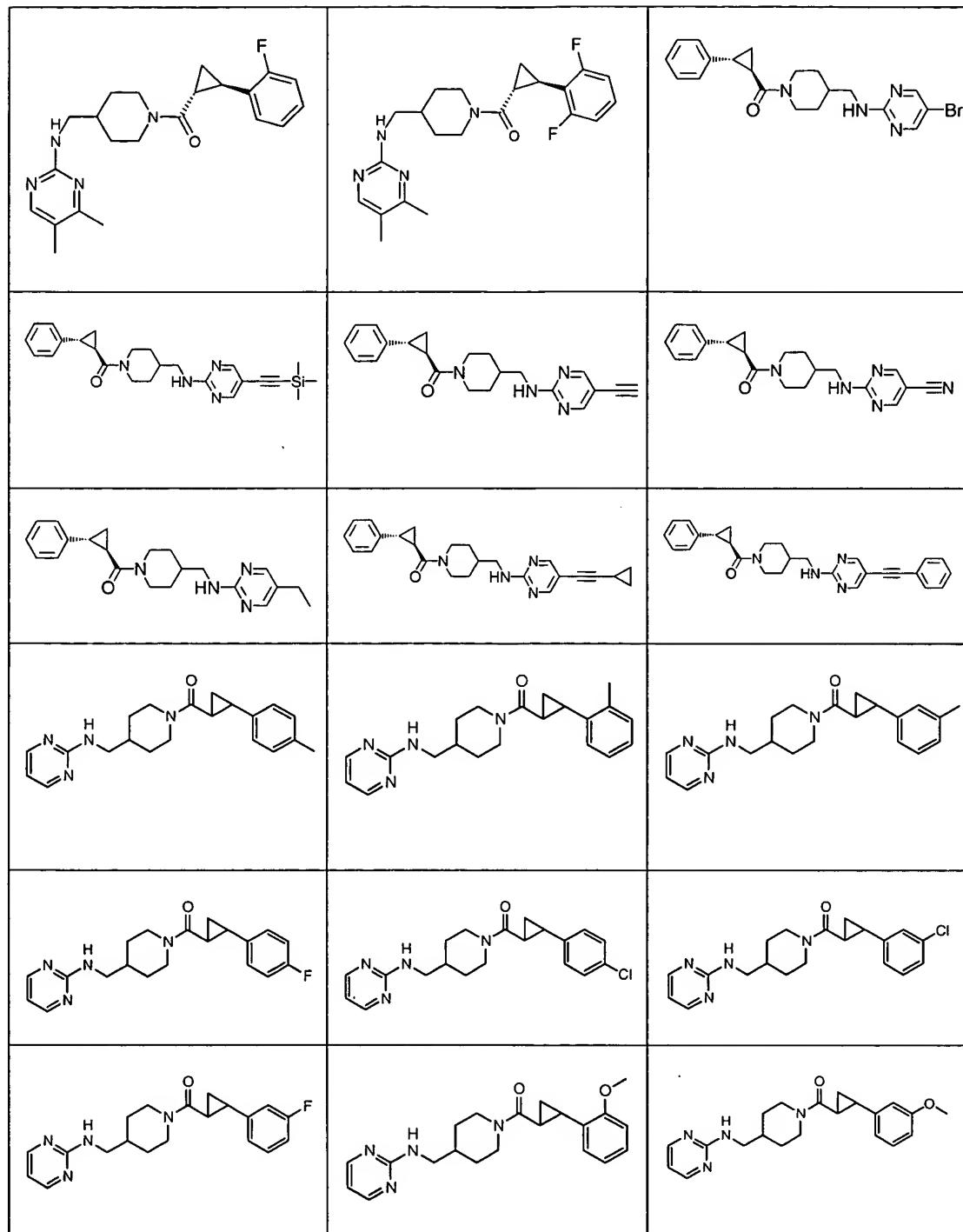


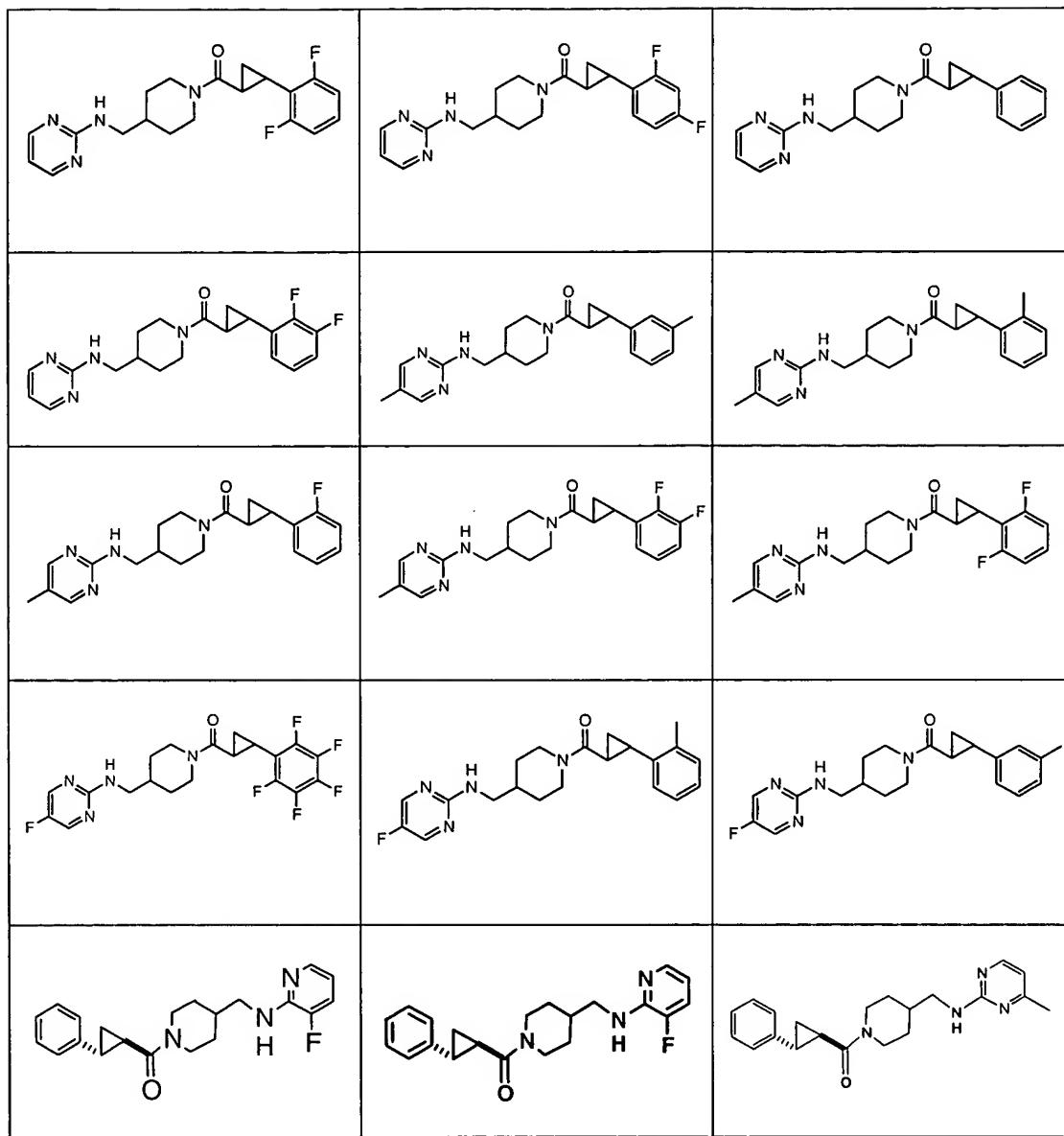
or a pharmaceutically acceptable salt thereof.

Claim 45(previously presented): The compound according to Claim 1, wherein said compound is



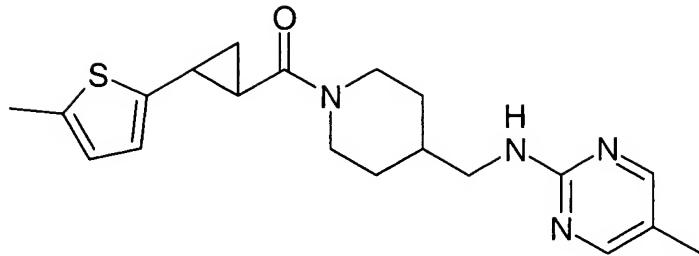






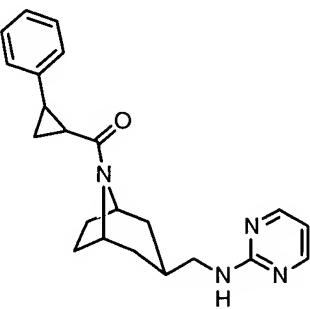
or a pharmaceutically acceptable salt thereof.

Claim 46(previously presented): The compound according to Claim 1, wherein said compound is



or a pharmaceutically acceptable salt thereof.

Claim 47(currently amended):  
A compound is represented by



or a pharmaceutically acceptable salt thereof.

Claim 48(original): A pharmaceutical composition comprising an inert carrier and an effective amount of a compound according to claim 1.

Claim 49(previously presented): A pharmaceutical composition comprising an inert carrier and an amount of a compound according to claim 1 effective to treat pain.

Claim 50(previously presented): A pharmaceutical composition comprising an inert carrier and an amount of a compound according to claim 1 effective to treat migraine, depression, anxiety, schizophrenia, Parkinson's disease, or stroke.

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Claim 51(original): A method of treating pain comprising a step of administering to one in need of such treatment an effective amount of a compound according to claim 1.

Claim 52(original): A method of treating migraine, depression, anxiety, schizophrenia, Parkinson's disease, or stroke comprising a step of administering to one in need of such treatment an effective amount of a compound according to claim 1.